AMENDMENTS TO THE CLAIMS

Please amend the claims as follows.

5 Claims 1-24 (canceled)

Claim 25 (withdrawn): A composition comprising 16α -bromo- 3β -hydroxy- 5α -androstan-17-one, 16α -bromo-2-oxa- 3β -hydroxy- 5α -androstan-17-one, 16α -bromo- 3β -hydroxy-11-oxa- 5α -androstan-17-one or 16α -bromo- 3β -hydroxy- 5α -androstan-17-one hemihydrate and one or more nonaqueous liquid excipients, wherein the composition comprises less than about 3% v/v water.

Claim 26 (withdrawn): The composition of claim 25 wherein the composition comprises less than about 0.3% v/v water.

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Claim 27 (withdrawn): The composition of claim 25 wherein the one or more nonaqueous liquid excipients are two or more of an alcohol, a polyethylene glycol, propylene glycol and benzyl benzoate.

Claim 28 (withdrawn): The composition of claim 25 wherein the composition is a parenteral formulation.

Claims 29-79 (canceled)

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Claim 80 (new): A method to treat or prevent an innate immune suppression condition in a human comprising administering to the human 1-10 mg/kg/day of a compound having the structure

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$$R^{1}$$
 R^{2}
 R^{2}
 R^{3}
 R^{4}
 R^{6}
 R^{3}
 R^{6}
 R^{3}
 R^{6}
 R^{3}
 R^{6}
 R^{3}

wherein, the dotted lines are optional double bonds and the hydrogen atom at the 5-position, if present, is in the α -configuration;

R¹ is -OR^{PR}, -SR^{PR}, an ester, a thioester, a phosphoester, a phosphothioester, a phosphonoester, a phosphiniester, an ether, a thioether, a carbonate or a thioacetal;

R² is -OR^{PR}, -SR^{PR}, =S, -CN, =NOH, =NOC(O)CH₃, an ester, a thioester, an ether, a thioether, an acyl group, a thioacyl group, a carbonate, a thioacetal, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl;

R³ is -H, -OR^{PR}, =O, -SR^{PR}, =S, -N(R^{PR})₂, -N₃, -CN, -NO₂, -F, -Cl, -Br, -I, an ester, a thioester, a thioacetal, an ether, a thioether, a carbamate, a carbonate, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl;

R⁴ is -OR^{PR}, -SR^{PR}, an ester, a thioester, phosphate, a phosphoester, a phosphothioester, a phosphonoester, a phosphiniester, a sulfite ester, a sulfate ester, an ether, a thioether, a carbonate, a thioacetal or a polymer;

R⁶ is -H, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl;

R⁹ is -CHR¹⁰- where R¹⁰ is -H, -OR^{PR}, =O, -SR^{PR}, =S, a halogen, an ester, an ether, a phosphoester, a carbonate, a thioacetal, a thioether, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl; and

R^{PR} independently are -H or an independently selected protecting group, whereby the number or activity of neutrophils in circulation in the human is increased.

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Claim 81 (new): The method of claim 80 wherein the innate immune suppression condition is associated with radiation, chemotherapy, aging, autologous bone marrow transplantation or stem cell transplantation.

Claim 82 (new): The method of claim 81 wherein the innate immune suppression condition is associated with the radiation or chemotherapy.

Claim 83 (new): The method of claim 82 wherein the compound has the structure

$$R^{1}$$
 R^{1}
 R^{2}
 R^{3}
 R^{4}
 R^{6}
 R^{6}
 R^{3}
 R^{4}
 R^{6}
 R^{3}
 R^{4}
 R^{6}
 R^{3}
 R^{4}
 R^{3}

Claim 84 (new): The method of claim 83 wherein R¹ is -OH, -SH, an ester, an ether or a carbonate.

Claim 85 (new): The method of claim 84 wherein R⁴ is -OH, -SH, an ester, phosphate, a phosphoester or an ether.

Claim 86 (new): The method of claim 85 wherein the compound has the structure

$$R^{9}$$
 R^{1}
 R^{2}
 R^{3}
 R^{4}
 R^{3}
 R^{4}
 R^{3}
 R^{4}
 R^{3}

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Claim 87 (new): The method of claim 86 wherein R³ is -F, -Cl, -Br, -I, -OH,=O, -SH, =S, an ester, an ether, a thioester, a thioacetal, a thioether, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl.

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Claim 88 (new): The method of claim 86 wherein R^9 is -CH₂-, -CH(OH)-, -C(O)-, or -CHR¹⁰-, wherein R^{10} is a halogen, an ester, an ether, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl.

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Claim 89 (new): The method of claim 82 wherein the compound has the structure

$$R^{1}$$
 R^{2}
 R^{2}
 R^{3}
 R^{4}
 R^{2}
 R^{3}
 R^{4}
 R^{2}
 R^{3}
 R^{4}
 R^{2}
 R^{3}

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and R² is -OH, an ester, an ether, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl.

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Claim 90. (new): The method of claim 82 wherein the compound is 3β ,17 β -dihydroxyandrost-5-ene, 3α ,17 β -dihydroxyandrost-5-ene, 16α -fluoro-17 β -dihydroxyandrost-5-ene, 16α -fluoro-17 α -dihydroxyandrost-5-ene, 16α -fluoro-17-oxoandrost-5-ene, 3β ,7 β ,17 β -trihydroxyandrost-5-ene, 3α ,7 β ,17 β -trihydroxyandrostane, 3β ,16 α ,17 β -trihydroxyandrostane or 3α ,16 α ,17 β -trihydroxyandrostane.

Claim 91 (new): The method of claim 90 wherein the compound is 3β ,17 β -

dihydroxyandrost-5-ene.

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Claim 92 (new): The method of claim 91 wherein the 3β ,17 β -dihydroxyandrost-5-ene is administered daily for 3, 4, 5, 6 or 7 consecutive days.

Claim 93 (new): The method of claim 91 wherein the 3β,17β-dihydroxyandrost-5-ene is parenterally administered for 5 consecutive days.

Claim 94 (new): The method of claim 93 wherein 50 mg, 75 mg, 100 mg, 150 mg, 200 mg, 250 mg or 300 mg per day of 3β ,17 β -dihydroxyandrost-5-ene is administered.

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Claim 95 (new): A method to treat or prevent an innate immune suppression condition in a non-human primate comprising administering to the non-human primate about 4-40 mg/kg/day of a compound having the structure

$$R^{6}$$
 R^{6}
 R^{6}
 R^{6}
 R^{6}
 R^{7}
 R^{7}
 R^{1}
 R^{2}
 R^{1}
 R^{2}
 R^{3}

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wherein, the dotted lines are optional double bonds and the hydrogen atom at the 5-position, if present, is in the α -configuration;

R¹ is -OR^{PR}, -SR^{PR}, an ester, a thioester, a phosphoester, a phosphothioester, a phosphonoester, a phosphiniester, an ether, a thioether, a carbonate or a thioacetal;

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R² is -OR^{PR}, =O, -SR^{PR}, =S, -CN, =NOH, =NOC(O)CH₃, an ester, a thioester, an ether, a thioether, an acyl group, a thioacyl group, a carbonate, a thioacetal, optionally substituted alkyl, optionally substituted alkynyl;

 R^3 is -H, -OR^{PR}, =O, -SR^{PR}, =S, -N(R^{PR})₂, -N₃, -CN, -NO₂, -F, -Cl, -Br, -I, an ester, a thioester, a thioacetal, an ether, a thioether, a carbonate, a

carbamate, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl;

R⁴ is -OR^{PR}, -SR^{PR}, an ester, a thioester, phosphate, a phosphoester, a phosphothioester, a phosphonoester, a phosphiniester, a sulfite ester, a sulfate ester, an ether, a thioether, a carbonate, a thioacetal or a polymer;

R⁶ is -H, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl;

R⁹ is -CHR¹⁰- where R¹⁰ is -H, -OR^{PR}, =O, -SR^{PR}, =S, a halogen, an ester, an ether, a phosphoester, a carbonate, a thioacetal, a thioether, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl; and

R^{PR} independently are -H or an independently selected protecting group, whereby the number or activity of neutrophils in circulation in the non-human primate is increased.

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Claim 96 (new): The method of claim 95 wherein the non-human primate is a cynomolgus monkey or a macaque monkey.

Claim 97 (new): The method of claim 96 wherein the innate immune suppression condition is associated with radiation, chemotherapy, autologous bone marrow transplantation or stem cell transplantation.

Claim 98 (new): The method of claim 97 wherein the innate immune suppression condition is associated with the radiation or chemotherapy.

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Claim 99 (new): The method of claim 98 wherein the compound has the structure

$$R^{9}$$
 R^{6}
 R^{7}
 R^{7}
 R^{7}

Claim 100 (new): The method of claim 99 wherein R¹ is -OH, -SH, an ester, an ether or a carbonate.

Claim 101 (new): The method of claim 100 wherein R⁴ is -OH, -SH, an ester, phosphate, a phosphoester or an ether.

Claim 102 (new): The method of claim 101 wherein the compound is 3β,17β-dihydroxyandrost-5-ene.

Claim 103 (new): The method of claim 101 wherein the 3β ,17 β -dihydroxyandrost-5-ene is administered daily for 3, 4, 5, 6 or 7 consecutive days.

15 Claim 104 (new): The method of claim 103 wherein the 3β ,17 β -dihydroxyandrost-5-ene is parenterally administered for 4, 5 or 6 consecutive days.

Claim 105 (new): The method of claim 104 wherein the 3β,17β-dihydroxyandrost-5-ene is administered for 5 consecutive days.

Claim 106 (new): A method to treat or prevent an innate immune suppression condition in a human, wherein the method comprises administering an effective amount of a compound for 3 to 15 consecutive days to the subject, wherein the compound has the structure

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$$R^{9}$$
 R^{2}
 R^{2}
 R^{3}
 R^{4}
 R^{4}
 R^{3}
 R^{4}
 R^{4}
 R^{4}
 R^{3}
 R^{4}
 R^{4}
 R^{4}
 R^{3}
 R^{4}
 R^{4}
 R^{4}
 R^{4}
 R^{4}
 R^{4}
 R^{4}

wherein, the dotted lines are optional double bonds and the hydrogen atom at the 5-position, if present, is in the α -configuration;

R¹ is -H, -OR^{PR}, -SR^{PR}, an ester, a thioester, a phosphoester, a phosphothioester, a phosphonoester, a phosphiniester, an ether, a thioether, a carbonate or a thioacetal;

R² is -OR^{PR}, =O, -SR^{PR}, =S, -CN, =NOH, =NOC(O)CH₃, an ester, a thioester, an ether, a thioether, an acyl group, a thioacyl group, a carbonate, a thioacetal, optionally substituted alkyl, optionally substituted alkynyl;

R³ is -H, -OR^{PR}, =O, -SR^{PR}, =S, -N(R^{PR})₂, -N₃, -CN, -NO₂, -F, -Cl, -Br, -I, an ester, a thioester, a thioacetal, an ether, a thioether, a carbamate, a carbonate, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl;

 R^4 in the β -configuration is $-OR^{PR}$, $-SR^{PR}$, an ester, a thioester, phosphate, a phosphoester, a phosphothioester, a phosphonoester, a phosphiniester, a sulfite ester, a sulfate ester, an ether, a thioether, a carbonate, a thioacetal, or a polymer;

 R^4 in the α -configuration is -H, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl;

R⁶ is -H, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl;

R⁹ is -CHR¹⁰- where R¹⁰ is -H, -OR^{PR}, =O, -SR^{PR}, =S, a halogen, an ester, an ether, a phosphoester, a carbonate, a thioacetal, a thioether, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl; and

R^{PR} independently are -H or an independently selected protecting group, whereby the numbers or activity of neutrophils in circulation in the human is increased.

Claim 107 (new): The method of claim 106 wherein the innate immune suppression condition is associated with radiation, chemotherapy, aging, autologous bone marrow transplantation or stem cell transplantation.

Claim 108 (new): The method of claim 107 wherein the innate immune suppression condition is associated with the radiation or chemotherapy.

Claim 109 (new): The method of claim 108 wherein the compound has the structure

$$R^{9}$$
 R^{6}
 R^{7}
 R^{8}

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Claim 110 (new): The method of claim 109 wherein R¹ is -OH, -SH, an ester, an ether or a carbonate.

Claim 111 (new): The method of claim 110 wherein R⁴ is -OH, -SH, an ester, phosphate, a phosphoester or an ether.

Claim 112 (new): The method of claim 111 wherein R³ is -F, -Cl, -Br, -I, -OH,=O, -SH, =S, an ester, an ether, a thioester, a thioacetal, a thioether, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl.

Claim 113 (new): The method of claim 110 wherein the compound is 3β ,17 β -dihydroxyandrost-5-ene, 3α ,17 β -dihydroxyandrost-5-ene, 16α -fluoro-17 β -dihydroxyandrost-5-ene, 3β ,7 β ,17 β -trihydroxyandrost-5-ene, 3α ,7 β ,17 β -trihydroxyandrostane, 3α ,16 β ,17 β -trihydroxyandrostane, 3α ,16 α ,17 β -trihydroxyandrostane or 3α ,16 α ,17 β -trihydroxyandrostane.

Claim 114 (new): The method of claim 113 wherein the compound is 3β,17β-dihydroxyandrost-5-ene.

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Claim 115 (new): The method of claim 114 wherein the 3β ,17 β -dihydroxyandrost-5-ene is administered daily for 4, 5 or 6 consecutive days.

Claim 116 (new): The method of claim 114 wherein the 3β ,17 β -dihydroxyandrost-5-ene is parenterally administered daily for 5 consecutive days.

Claim 117 (new): The method of claim 116 wherein about 1.0 mg/kg/day, about 1.5 mg/kg/day, about 2 mg/kg/day, about 2.5 mg/kg/day, about 3.0 mg/kg/day, about 4 mg/kg/day or about 6 mg/kg/day of the 3β ,17 β -dihydroxyandrost-5-ene is administered.

Claim 118 (new): The method of claim 116 wherein 50 mg, 75 mg, 100 mg, 150 mg, 200 mg, 250 mg or 300 mg of the 3β ,17 β -dihydroxyandrost-5-ene is

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administered each day.